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(presently amended) A method for modifying a carbohydrate, comprising the steps of:

- (a) selecting providing at least one <u>purified</u> glycosidase of <u>defined</u> substrate specificity obtainable from <u>Xanthomonas holcicola</u>, <u>Xanthomonas manihotis</u>, or <u>Xanthomonas oryzae</u> wherein the glycosidase is selected from the group consisting of a fucosidase, a mannosidase, a xylosidase, a glucosidase, a galactosidase, N-acetylglucosaminidase and a hexosaminidase:
- (b) cleaving a selected glycosidic bond between constituent monosaccharides of the carbohydrate by means of the glycosidase; and
  - (c) forming a modified carbohydrate.
  - 8 (canceled)
  - 9.(canceled)
- 10. (previously presented) The method according to claim 7, wherein the-modified carbohydrate has altered immunogenic properties compared with the carbohydrate prior to modification.
- 11. (previously presented) The method according to claim 7, wherein step (b) further comprises cleaving Fuc $\alpha$ 1-2R linkage.
  - 12 cancelled
  - cancelled

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14 (new) A method according to claim 7, wherein the glycosidase is selected from the group consisting of a fucosidase, a mannosidase, a xylosidase, a glucosidase, a galactosidase, N-acetylglucosaminidase and a hexosaminidase.

- 15. (new) A method according to claim 7, wherein the glycosidase is selected from a  $\beta$ 1-3>>4 galactosidase, an  $\alpha$ -1-2,3 mannosidase, a  $\beta$ -glucosidase, an  $\alpha$ 1-3,4 fucosidase, an  $\alpha$ 1-2 fucosidase, a  $\beta$ -N-acetylglucosaminidase,  $\beta$ -GlcNAc, an  $\alpha$  1-6 mannosidase, an  $\alpha$ -1-3,6 galactosidase, an  $\alpha$ -1-3,6 mannosidase, a  $\beta$ -xylosidase and a  $\beta$ -mannosidase.
- 16. (new) A method according to claim 7, wherein step (a) further comprises determining the defined substrate specificity using a fluorescent chromophore.
- 17. (new) A method according to claim 16, wherein the fluorescent chromophore is 7-aminocoumarin.
- 18. (new) A method according to claim 7, wherein step (b) further comprises measuring a hydrolysis product resulting from cleavage of the glycosidic bond using thin layer silica gel chromatography.